IN THE CLAIMS

Please cancel claims 1, 2, 8, 10, 16-17, 19, 22, 25-26 and 28 without prejudice or disclaimer.

Please amend claims 3, 9, 13, 15 and 28 as follows.

1-2. (Canceled)

- 3. (Currently Amended): An isolated polynucleotide encoding [a polypeptide of claim 1] an isolated polypeptide comprising an amino acid sequence selected from the group consisting of:
 - a) the amino acid sequence of SEQ ID NO:8, and
 - b) a naturally occurring amino acid sequence having at least 90% sequence identity to the amino acid sequence of SEQ ID NO:8.
 - 4. (Original): An isolated polynucleotide encoding a polypeptide of claim 2.
- 5. (Currently Amended): An isolated polynucleotide of claim 4 [selected from the group consisting of SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:27, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:32, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:36, SEQ ID NO:37, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41, SEQ ID NO:42, and SEQ ID NO:44] comprising SEQ ID NO:30.
- 6. (Original): A recombinant polynucleotide comprising a promoter sequence operably linked to a polynucleotide of claim 3.
 - 7. (Original): A cell transformed with a recombinant polynucleotide of claim 6.
 - 8. (Canceled)

- 9. (Currently Amended): A method for producing [a polypeptide of claim 1] an isolated polypeptide, said polypetide comprising an amino acid sequence selected from the group consisting of:
 - a) the amino acid sequence of SEQ ID NO:8, and
 - b) a naturally occurring amino acid sequence having at least 90% sequence identity to the amino acid sequence of SEQ ID NO:8,

the method comprising:

[a] <u>i</u>) culturing a cell under conditions suitable for expression of the polypeptide, wherein said cell is transformed with a recombinant polynucleotide, and said recombinant polynucleotide comprises a promoter sequence operably linked to a polynucleotide encoding [the polypeptide of claim 1] <u>the polypeptide</u>,

and

- [b] <u>ii</u>) recovering the polypeptide so expressed.
- 10. (Canceled)
- 11. (Currently Amended): An isolated polynucleotide comprising a polynucleotide sequence selected from the group consisting of:
 - a) a polynucleotide sequence [selected from the group consisting of SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:27, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:32, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:36, SEQ ID NO:37, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41, SEQ ID NO:42, and SEQ ID NO:44] comprising SEQ ID NO:30,
 - a naturally occurring polynucleotide sequence having at least 70% sequence identity to a polynucleotide sequence [selected from the group consisting of SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:27, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:32, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:36, SEQ ID NO:37, SEQ

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ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41, SEQ ID NO:42, and SEQ ID NO:44] comprising SEQ ID NO:30,

- c) a polynucleotide sequence complementary to a),
- d) a polynucleotide sequence complementary to b), and
- e) an RNA equivalent of a)-d).
- 13. (Original): A method for detecting a target polynucleotide in a sample, said target polynucleotide having a sequence of a polynucleotide of claim 11, the method comprising:
- a) hybridizing the sample with a probe comprising at least 20 contiguous nucleotides comprising a sequence complementary to said target polynucleotide in the sample, and which probe specifically hybridizes to said target polynucleotide, under conditions whereby a hybridization complex is formed between said probe and said target polynucleotide or fragments thereof, and
- b) detecting the presence or absence of said hybridization complex, and, optionally, if present, the amount thereof.
- 15. (Original): A method for detecting a target polynucleotide in a sample, said target polynucleotide having a sequence of a polynucleotide of claim 11, the method comprising:
- a) amplifying said target polynucleotide or fragment thereof using polymerase chain reaction amplification, and
- b) detecting the presence or absence of said amplified target polynucleotide or fragment thereof, and, optionally, if present, the amount thereof.

16-17, 19, 22, 25-26. (Canceled)

- 28. (Original)A method for assessing toxicity of a test compound, said method comprising:
 - a) treating a biological sample containing nucleic acids with the test compound;
- b) hybridizing the nucleic acids of the treated biological sample with a probe comprising at least 20 contiguous nucleotides of a polynucleotide of claim 11 under conditions whereby a

specific hybridization complex is formed between said probe and a target polynucleotide in the biological sample, said target polynucleotide comprising a polynucleotide sequence of a polynucleotide of claim 11 or fragment thereof;

- c) quantifying the amount of hybridization complex; and
- d) comparing the amount of hybridization complex in the treated biological sample with the amount of hybridization complex in an untreated biological sample, wherein a difference in the amount of hybridization complex in the treated biological sample is indicative of toxicity of the test compound.